

**CLAIMS**

What is claimed:

1. A method of treating breast cancer which comprises administering, to a subject suffering from breast cancer,

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a first amount of anti-estrogenic steroid agent, effective to reduce the level or activity of at least one estrogenic steroid in the subject, and

10 a second amount of an immunological agent, effective to contribute to the development of a protective immune response to said breast cancer,

15 where said first and second amounts are, at least in combination, therapeutically effective against at least some breast cancers.

2. The method of claim 1 where said agents are administered concurrently.

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3. The method of claims 1 or 2 where said immunological agent comprises at least one immunogen, said immunogen comprising at least one breast cancer-associated epitope.

4. The method of claim 3 where at least one epitope is a MUC1 epitope.

5. The method of claim 3 where at least one epitope is a 5 carbohydrate epitope.

6. The method of claim 3 in which said immunogen comprises STn.

10 7. The method of claim 6 in which said immunogen comprising STn is an STn-KLH conjugate.

8. The method of claim 7 in which the conjugate is an aggregated conjugate.

15 9. The method of claim 7 or 8 in which the conjugate has a NANA content of about 7%.

10. The method of any one of claims 1-9 in which the 20 anti-estrogenic steroid agent comprises at least one antiestrogen.

11. The method of claim 10 in which at least one antiestrogen is a steroidal antiestrogen.

12. The method of claim 1 in which at least one anti-estrogenic steroid agent is fulvestrant.

13. The method of claim 10 in which at least one  
5 antiestrogen is a nonsteroidal antiestrogen.

14. The method of claim 13 in which at least one  
nonsteroidal antiestrogen is selected from the group  
consisting of toremifene, tamoxifen, droloxifene and  
10 trioxifene

15. The method of any one of claims 1-14 in which the  
anti-estrogenic steroid agent comprises at least one  
aromatase inhibitor.

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16. The method of claim 15 in which at least one  
aromatase inhibitor is selected from the group consisting  
of aminoglutethimide, anastrozole, vorozole, letrozole,  
lairozole, megastrole, exemestane and formestane.

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17. The method of any one of claims 1-16, further  
comprising administration of at least one progestin which  
protects against breast cancer.

18. The method of claim 17 in which at least one progestin is progesterone.

19. The method of any one of claims 1-18, further comprising administration of at least one anti-progestin which protects against breast cancer.

20. The method of any one of claims 1-19 in which the anti-estrogenic steroid agent comprises geoselin acetate or megestrol acetate.

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21. The method of any one of claims 1-20 in which the combination of the anti-estrogenic steroid agent and the immunological agent is synergistically effective against breast cancer.

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22. The method of any one of claims 1-21, further comprising administration of a therapeutically effective amount of at least one chemotherapeutic agent other than an anti-estrogenic steroid agent.

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23. The method of claim 22 in which at least one chemotherapeutic agent is an anthracycline.

24. The method of claim 23 in which at least one anthracycline is selected from the group consisting of doxorubicin, daunorubicin, epirubicin, and idarubicin.

5 25. The method of claim 22 in which at least one chemotherapeutic agent is a taxane.

26. The method of claim 25 in which at least one taxane is paclitaxel or docetaxel.

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27. The method of any one of claims 1-26 in which the anti-estrogenic steroid agent comprises at least one compound which antagonizes at least one estrogen receptor by competitively inhibiting the binding of an estrogen to 15 that receptor without itself activating that receptor.

28. The method of claim 27 in which said receptor antagonist is not an agonist for any estrogen receptor.

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29. The method of claim 27 in which said receptor inhibitor is also an agonist of at least one other estrogen receptor, and consequently is a SERM.

25 30. The method of claim 29 in which said SERM is selected from the group consisting of tamoxifen, toremifene,

droloxifen, clomifene, arzoxifene, raloxifene, LY 117018  
and SERM EM-652.

31. The method of any one of claims 1-30 in which the  
5 breast cancer is a metastatic breast cancer.

32. A therapeutic composition comprising (a) at least one  
anti-estrogenic steroid agent, and (b) at least one  
immunogenic agent, which, when administered according to  
10 a suitable therapeutic schedule, is therapeutically  
effective against breast cancer.

33. A kit comprising a first container comprising at  
least one dose of at least one anti-estrogenic steroid  
15 agent, and a second container comprising at least one  
dose of at least one immunogenic agent, where said agents  
are, at least in combination, therapeutically effective  
against breast cancer.

20 34. Use of (1) a first amount of anti-estrogenic steroid  
agent, effective to reduce the level or activity of at  
least one estrogenic steroid in the subject, and (2)  
a second amount of an immunological agent, effective to  
contribute to the development of a protective immune  
25 response to breast cancer,

where said first and second amounts are, at least in combination, therapeutically effective against at least some breast cancers,

5 in the manufacture of one or more compositions for the treatment of breast cancer.

35. Use of an immunological agent, effective to contribute to the development of a protective immune 10 response to breast cancer, in a manufacture of a composition for the treatment of breast cancer in a subject who is receiving or has received treatment with an anti-estrogenic steroid agent, effective to reduce the level or activity of at least one estrogenic steroid in 15 the subject.

36. Use of an anti-estrogenic steroid agent, effective to reduce the level or activity of at least one estrogenic steroid in the subject, in the manufacture of a 20 composition for the treatment of breast cancer in a subject who is receiving or has received treatment with an immunological agent, effective to contribute to the development of a protective immune response to said breast cancer.

37. The use of claim 34, wherein the agent is or agents are as set forth in any of claims 2-30.

38. The composition of claim 32, wherein the agent is or  
5 agents are as set forth in any of claims 2-30.

39. The kit of claim 33, wherein the agent is or agents are as set forth in any of claims 2-30.